Double-sided back muscle numbing for pain relief pain after having the gallbladder removed using keyhole surgery

Submission date	Recruitment status	Prospectively registered
16/09/2023	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
21/09/2023	Completed	Results
Last Edited	Condition category	Individual participant data
21/09/2023	Surgery	Record updated in last year

Plain English summary of protocol

Background and study aims

Laparoscopic cholecystectomy is a specific type of surgery where the gallbladder is removed using small incisions and a camera. An erector spinae block is a medical procedure in which a local anesthetic (a medication that numbs sensation) is injected near the erector spinae muscles along the spine. This is done to block or numb the nerves that transmit pain signals from the surgical area to the brain. The purpose of using this block is to provide effective pain relief during and after the surgery. By numbing the nerves in the area, it helps reduce the amount of pain the patient feels, making the surgical procedure more comfortable and improving the postoperative recovery experience. It's a part of the overall pain management strategy for the surgery, potentially leading to better pain control and a smoother recovery process for the patient.

Who can participate?

Patients getting laparoscopic cholecystectomy

What does the study involve?

Patients were split into two groups: one receiving general anesthesia plus regular pain control, and the other receiving both of those plus an erector spinae block.

What are the possible benefits and risks of participating?

The potential advantages include better pain relief, while the risks involve the use of ultrasound to guide the block.

Where is the study run from? Universidad del Sinu (Colombia)

When is the study starting and how long is it expected to run for? May 2021 to February 2023

Who is funding the study? Universidad del Sinu (Colombia)

Who is the main contact? Mauro Causil, jjmoralesb@unal.edu.co

Contact information

Type(s)

Principal Investigator

Contact name

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

08012021

Study information

Scientific Title

Bilateral erector spinae block for postoperative analgesia in laparoscopic cholecystectomy: A randomized clinical trial

Study objectives

ESPB improves postoperative analgesia and reduces opioid consumption

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 08/01/2021, Clinica La Concepcion Ethics Comittee (CALLE 38 52 249 LOS ALPES, Sincelejo, 700001, Colombia; (+57) 5 2749180; info@laconcepcion.org), ref: 08012021

Study design

Prospective interventional randomized clinical trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet.

Health condition(s) or problem(s) studied

Improvement of analgesia in patients undergoing laparoscopic cholecystectomy

Interventions

There were 2 study arms: one was general anaesthesia + standard analgesia and the other arm was general anaesthesia + standard analgesia + bilateral US-guided erector spinae block.Follow up was 48 hrs after surgery to record pain scores. The randomization process was done using Epi Dat 4.2.

Intervention Type

Other

Primary outcome measure

Pain measured using a 0 (least)-10 (worst) pain score 1, 6, 12, 24 and 48 hr after surgery

Secondary outcome measures

Requirement of opioid (Yes or No) 1 hour after surgery

Overall study start date

01/05/2021

Completion date

28/02/2023

Eligibility

Key inclusion criteria

- 1. More than 18 years old
- 2. ASA I,II
- 3. Undergoing laparoscopic cholecystectomy

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Upper age limit

100 Years

Sex

Both

Target number of participants

68

Total final enrolment

68

Key exclusion criteria

- 1. IMC >30
- 2. Open cholecystectomy
- 3. Allergy to local anesthetics
- 4. Recent opioid use

Date of first enrolment

01/08/2021

Date of final enrolment

28/02/2023

Locations

Countries of recruitment

Colombia

Study participating centre Clinica La Concepcion

CALLE 38 52 249 LOS ALPES Sincelejo Colombia 700001

Sponsor information

Organisation

Universidad del Sinú

Sponsor details

Calle 31 N° 3-06 Montería Colombia 230001 +57 7811717 admisiones@unisinu.edu.co

Sponsor type

University/education

Website

https://www.unisinu.edu.co

ROR

https://ror.org/013ys5k90

Funder(s)

Funder type

University/education

Funder Name

Universidad del Sinú

Results and Publications

Publication and dissemination plan

Planned publication is a high impact peer reviewed journal

Intention to publish date

18/03/2024

Individual participant data (IPD) sharing plan

The datasets generated during and or analyses during the current study will be available upon request from jjmoralesb@unal.edu.co

IPD sharing plan summary Available on request